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Selective Liquid Phase Hydrogenation of Benzaldehyde to Benzyl Alcohol Over Alumina Supported Gold

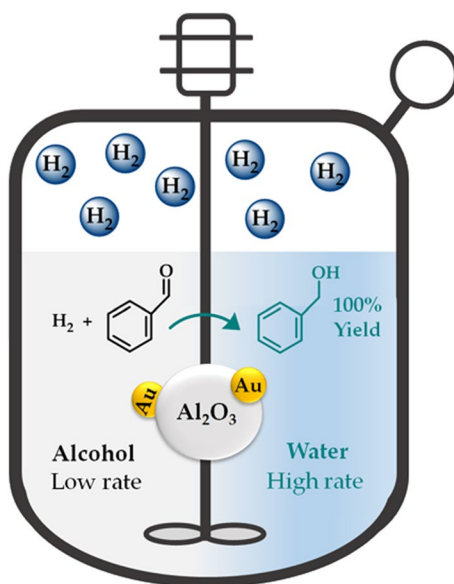
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Abstract

We report for the first time 100% benzyl alcohol yield from the liquid phase ($T=353\text{ K}$, $P=9\text{ bar}$) hydrogenation of benzaldehyde over $\text{Au}/\text{Al}_2\text{O}_3$. Under the same reaction conditions, a benchmark $\text{Pt}/\text{Al}_2\text{O}_3$ catalyst promoted the formation of toluene and benzene as hydrogenolysis by-products. Reaction kinetics was subjected to a Hammett treatment and the reaction constant ($\rho=0.9$) was found to be consistent with a nucleophilic mechanism. A solvent (alcohol, water and alcohol + water) effect is demonstrated and ascribed to competitive adsorption where solvation by polar (water) facilitates benzaldehyde activation.

Graphic Abstract



Keywords Selective hydrogenation · Benzaldehyde · Benzyl alcohol · Water as solvent · $\text{Au}/\text{Al}_2\text{O}_3$ · $\text{Pt}/\text{Al}_2\text{O}_3$

1 Introduction

The production of benzyl alcohol by the selective reduction of benzaldehyde is an important commercial process with multiple applications as solvent for inks, paints and lacquers [1]. Oxide supported Ru [2], Pd [3, 4], Ni [5, 6] Cu [7] and Pt [8, 9] catalysts have been employed, typically in batch liquid phase at high pressure (20–40 bar) [1, 2, 5, 8, 10,

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[11]. The target of 100% benzyl alcohol yield (Scheme 1, path I) remains a challenge as undesired hydrogenolysis (to toluene and/or benzene, paths II–V [12]) is difficult to circumvent with reported alcohol yields in the 40–75% range [2–6, 9, 13]. Application of Au in hydrogenation, though less developed than use in oxidation, has been the focus of recent research [14–19]. In their review, Hari and Yaakob [20] established the potential of Au for selective liquid phase hydrogenation with enhanced alcohol selectivity in the conversion of aldehydes such as crotonaldehyde [21] and citral [22]. We provide here the first reported application of (Al₂O₃) supported Au in the liquid phase hydrogenation of benzaldehyde.

Gold delivers lower activity when compared with conventional transition metal catalysts due to a less effective activation/dissociation of H₂ [23, 24]. Hydrogenation activity in liquid phase operation can be influenced by the solvent [25]. This has been related to differences in dielectric constant (ϵ) [26–28], bonding capacity (α) [28, 29] and H₂ solubility [26]. Aramendía et al. [30] and Bertero et al. [27] found that the rate of acetophenone hydrogenation decreased with increasing ϵ of C₁–C₃ alcohols, which they ascribed to solvation effects that influenced reactant adsorption. On the other hand, Wan et al. [28] observed an increased hydrogenation rate with increasing α , linked to interaction between

protic solvents and 2-butanone by hydrogen bonding that lowered the activation energy barrier. Drelinkiewicz et al. [26] concluded that acetophenone hydrogenation was more influenced by solvent polarity than H₂ solubility. Organic solvents such as ethanol [3, 4, 9, 10, 13], dioxane [5] and alkanes (*n*-octane [3] and dodecane [31]) have been used in the liquid phase hydrogenation of benzaldehyde. Water as a benign green reaction medium has not been considered to any great extent. It is nonetheless worth flagging the work of Wang and co-workers [32] who demonstrated higher activity in the chemoselective hydrogenation of α , β -unsaturated carbonyl compounds over Au/CeO₂ in water relative to organic solvents (ethanol, isopropanol, dioxane and cyclohexane).

In this work, we examine the role of solvent (*n*-pentanol, *n*-butanol, ethanol, H₂O and ethanol + H₂O mixtures) in determining benzaldehyde hydrogenation over Au/Al₂O₃. We have targeted 100% benzyl alcohol yield as an objective. The effect of *para*-positioned substituents (–OCH₃, –CH₃, –CH₂CN and –CN) on rate has also been considered. Given the established application of supported Pt in benzaldehyde hydrogenation [8, 10, 11], we employed a commercial Pt/Al₂O₃ as a suitable benchmark catalyst.

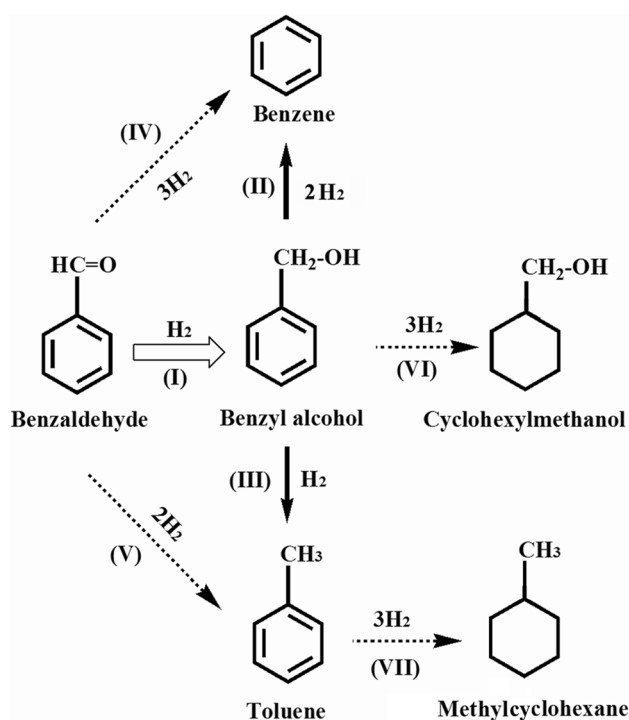
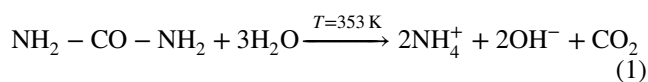
2 Experimental

2.1 Materials

The Al₂O₃ support and (0.7% w/w) Pt/Al₂O₃ were purchased from Puralox (Condea Vista. Co.) and Alfa Aesar, respectively. The gold precursor (HAuCl₄; 99.999%, Sigma-Aldrich) and urea ($\geq 99\%$; Riedel-de H  en) were used as received. All the gases used (H₂, N₂, O₂ and He) were of high purity (99.9%, BOC gases). The reactants (*p*-methoxybenzaldehyde, *p*-tolualdehyde, benzaldehyde, benzyl alcohol and *p*-cyanobenzaldehyde; $\geq 98\%$, Aldrich; *p*-formylphenyl acetonitrile, 96.5%, Apollo Scientific) and solvents (ethanol, butanol, pentanol; $\geq 99.5\%$, Aldrich) were used as supplied without further purification.

2.2 Catalyst Preparation and Activation

A 1% w/w Au/Al₂O₃ was prepared by deposition–precipitation using urea as basification agent. An aqueous mixture of urea (tenfold excess) and HAuCl₄ in water (100 cm³, 25×10^{-3} g cm^{−3}, pH = 2) was added to the support (ca. 10 g). The suspension was stirred and heated to 353 K (at 2 K min^{−1}) and the pH progressively increased to ca. 7 after 3 h as a result of thermally induced urea decomposition.



Scheme 1 Reaction pathway associated with benzaldehyde hydrogenation; pathway to the target alcohol (open arrow), by-products detected in this work (solid arrows) and reported in the literature [12] (dashed arrows)

The solid obtained was separated by centrifugation, washed with deionised water (with centrifugation between each washing) and dried in He ($45 \text{ cm}^3 \text{ min}^{-1}$) at 2 K min^{-1} to 373 K , which was maintained for 5 h . Au/ Al_2O_3 and Pt/ Al_2O_3 were crushed and sieved to $75 \mu\text{m}$ average particle diameter (ATM fine test sieves) and activated in $60 \text{ cm}^3 \text{ min}^{-1} \text{ H}_2$ at 2 K min^{-1} to 523 K (Au/ Al_2O_3 [33]) and 773 K (Pt/ Al_2O_3) [34]. Post activation, samples were cooled to 298 K and passivated in $1\% \text{ v/v O}_2/\text{He}$ for ex situ analysis.

2.3 Catalyst Characterisation

The metal (Au and Pt) content was measured by inductively coupled plasma-optical emission spectrometry (ICP-OES, Vista-PRO, Varian Inc.) from the (fivefold) diluted extract in HF (10 cm^3 , 0.1 M). Catalyst activation and chemisorption (at 353 K) measurements were conducted using the commercial CHEM-BET 3000 (Quantachrome) unit. The sample was loaded into a U-shaped quartz cell (3.76 mm i.d.), heated in $17 \text{ cm}^3 \text{ min}^{-1}$ (Brooks mass flow controlled), $5\% \text{ v/v H}_2/\text{N}_2$ at 2 K min^{-1} to 523 – 773 K and maintained until the signal returned to baseline. Samples were swept with $65 \text{ cm}^3 \text{ min}^{-1} \text{ N}_2$ for 1.5 h , cooled to 353 K and subjected to H_2 chemisorption using a pulse (10 – $50 \mu\text{l}$) titration procedure. Hydrogen pulse introduction was repeated until the signal area was constant, indicating surface saturation. Metal particle morphology (size and shape) was determined by scanning transmission (JEOL 2200FS field emission gun-equipped TEM unit) electron microscopy (STEM), employing Gatan DigitalMicrograph 1.82 for data acquisition/manipulation. Samples were dispersed in acetone and deposited on a holey carbon/Cu grid (300 Mesh). Up to 1000 individual particles were counted and the surface area mean metal (Au, Pt) diameter (d_{STEM}) calculated from

$$d_{\text{STEM}} = \frac{\sum_i n_i d_i}{\sum_i n_i} \quad (2)$$

where n_i is the number of particles of diameter d_i .

2.4 Catalytic System

The liquid phase hydrogenation reactions ($T = 353 \text{ K}$; $P = 9.0 \text{ bar}$) were carried out in a commercial batch stirred stainless steel reactor (100 cm^3 autoclave, Parr reactor) equipped with a H_2 supply system (GCE-Druva). The temperature was maintained at $353 \pm 1 \text{ K}$ using a process controller (Scientific & Medicine Products Ltd). At the beginning of each run, the aldehyde (or benzyl alcohol) solution (40 cm^3 of pentanol, butanol, ethanol, H_2O or ethanol + H_2O ; 0.05 M) and catalyst were charged and flushed three times

with N_2 . The system was heated to the reaction temperature, pressurised and the stirring engaged (time $t = 0$ for reaction). In a series of blank tests, there was negligible conversion in the absence of catalyst or without H_2 . The initial molar reactant to metal ratio spanned the range 2×10^3 – 11×10^3 . A liquid sampling system via syringe with in-line filters allowed a controlled withdrawal of aliquots ($\leq 1.0 \text{ cm}^3$) from the reactor. The concentration of reactant/product was determined from the total mass balance in the reaction mixture. Fractional conversion of benzaldehyde ($X_{\text{Benzaldehyde}}$) is defined as

$$X_{\text{Benzaldehyde}} (-) = \frac{C_{\text{Benzaldehyde},0} - C_{\text{Benzaldehyde}}}{C_{\text{Benzaldehyde},0}} \quad (3)$$

where subindex ‘0’ refers to initial concentration. Benzaldehyde consumption rate (R , $\text{mol mol}_{\text{metal}}^{-1} \text{ h}^{-1}$) was determined from a linear regression of the temporal benzaldehyde concentration profiles at $X_{\text{Benzaldehyde}} < 0.35$ [35] according to:

$$R (\text{mol mol}_{\text{metal}}^{-1} \text{ h}^{-1}) = \left(\frac{\Delta C_{\text{Benzaldehyde}}}{\Delta t} \right) \cdot \left(\frac{1}{n_{\text{metal}}} \right) \quad (4)$$

where n_{metal} is the metal concentration ($\text{mol}_{\text{metal}} \text{ cm}^{-3}$). Turnover frequency (TOF , s^{-1}) was calculated using metal dispersion measurements from STEM as described elsewhere [36]. Selectivity to benzyl alcohol ($S_{\text{Benzyl alcohol}}$) is given by

$$S_{\text{Benzyl alcohol}} (\%) = \frac{C_{\text{Benzyl alcohol}}}{C_{\text{Benzaldehyde},0} - C_{\text{Benzaldehyde}}} \times 100 \quad (5)$$

Product composition was determined by gas chromatography using a Perkin-Elmer AutoSystem XL chromatograph equipped with a programmed split/splitless injector and a flame ionisation detector, employing a DB-1 capillary column ($i.d. = 0.33 \text{ mm}$, length = 50 m , film thickness = $0.20 \mu\text{m}$). Data acquisition and manipulation were performed using the TotalChrom Workstation (Version 6.3.2 for Windows) chromatography data system. Repeated reaction runs with the same batch of catalyst delivered raw data reproducibility and carbon mass balances within $\pm 5\%$.

3 Results and Discussion

3.1 Catalyst Characterisation

The metal loading, particle size (from STEM analysis) and H_2 chemisorption values are given in Table 1. Both supported catalysts exhibit similar loading where the metal phase takes the form of pseudo-spherical particles (Fig. 1I) over the 1 – 8 nm size range (Fig. 1II). This equivalence permits a direct comparison of catalytic response. Hydrogen uptake on supported Au is sensitive to metal size and

Table 1 Metal content, mean metal particle size from STEM analysis (d_{STEM}), H_2 chemisorption (at 353 K) and catalytic activity expressed as reaction rate (R) and turnover frequency (TOF) in the liquid phase hydrogenation of benzaldehyde (using water as solvent): $T=353\text{ K}$; $P=9\text{ bar}$

	Au/ Al_2O_3	Pt/ Al_2O_3
Metal content w/w (%)	1.1	0.7
d_{STEM} (nm)	3.6	1.7
H_2 uptake ($\mu\text{mol g}^{-1}$)	2.8	22.7
R ($\text{mol mol}_{\text{metal}}^{-1} \text{h}^{-1}$)	78	446
$\text{TOF} \times 10^{-3}$ (s^{-1})	65	186

dissociative adsorption is favoured on smaller (1–10 nm) particles [37]. The eightfold lower H_2 uptake on Au/ Al_2O_3 relative to Pt/ Al_2O_3 (Table 1) can be linked to the high activation barrier for H_2 adsorption on Au [38] and greater capacity of Pt to dissociate H_2 [39, 40].

3.2 Liquid Phase Hydrogenation of Benzaldehyde

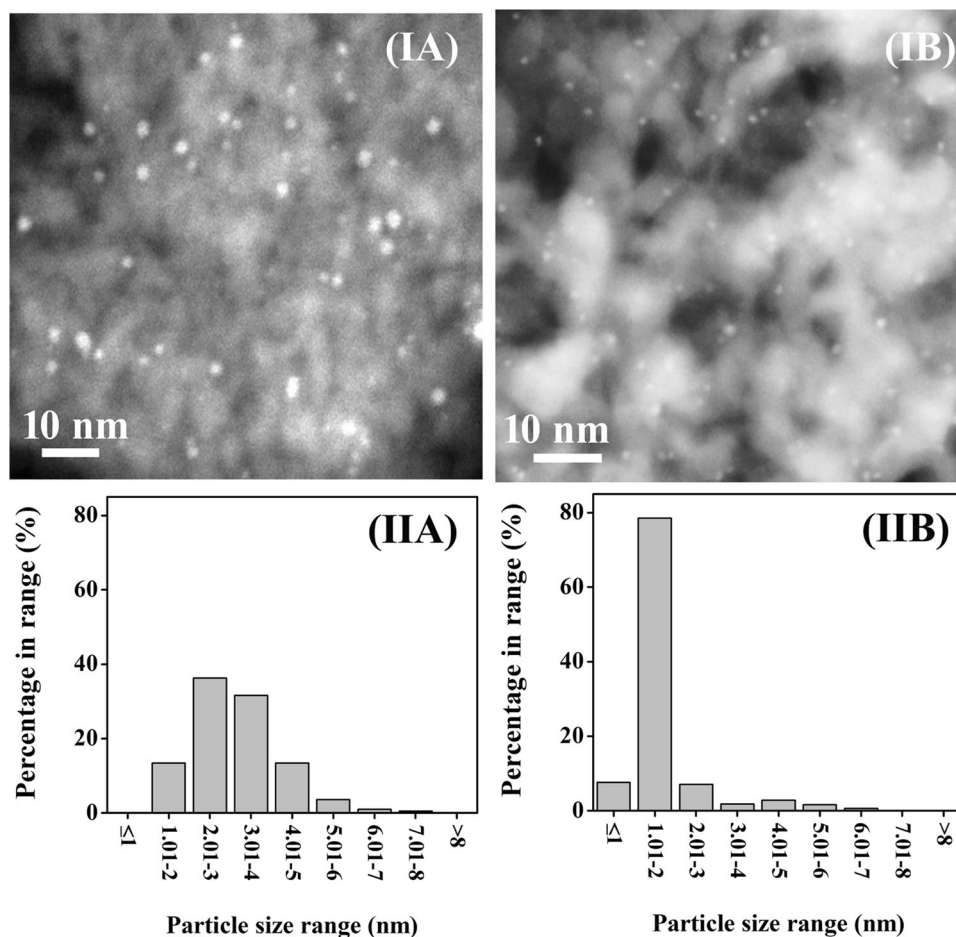
Reaction selectivity is challenging in benzaldehyde hydrogenation [1–6, 9, 13] with possible by-products, as shown in

Scheme 1. Step I represents the target carbonyl group reduction. Subsequent hydrogenolysis results in the formation of benzene (step II) and toluene (step III) [41, 42]. Benzene and toluene can also be formed directly from benzaldehyde (steps IV and V) [41]. Further hydrogenation of benzyl alcohol to cyclohexylmethanol (step VI) has been reported for reaction over Ru/C [43] with toluene reduction to methylcyclohexane (step VII) over Ni/ Al_2O_3 [6]. Under all reaction conditions used in this study, benzaldehyde was converted solely to the target benzyl alcohol over Au/ Al_2O_3 .

3.2.1 Reaction Under Kinetic Control

Benzaldehyde hydrogenation rate was determined from the linear variation of concentration with time, as shown in Fig. 2I. We carried out a series of tests in order to minimise diffusion and mass transfer limitations and ensure the reaction was operated under kinetic control. The effect of variations in agitation speed (Fig. 2IA) and catalyst mass (Fig. 2IB) on reaction rate was examined as two well-established diagnostic tests to assess external and internal transfer constraints [44]. An increase in stirring speed from 300 to 700 rpm was accompanied by a proportional increase in rate.

Fig. 1 I Representative STEM images with II associated particle size distribution for (A) Au/ Al_2O_3 and (B) Pt/ Al_2O_3



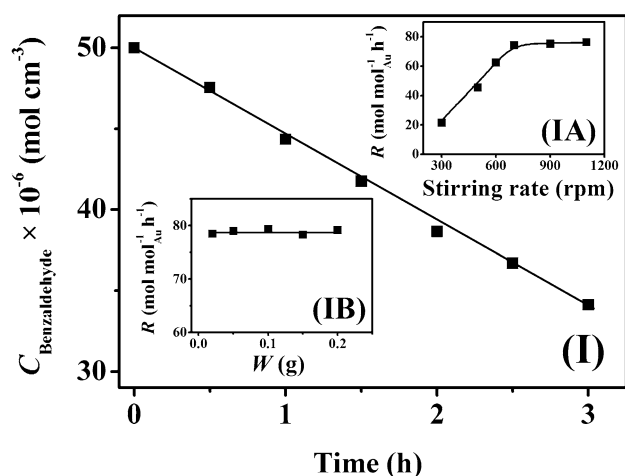


Fig. 2 I Variation of benzaldehyde concentration in water ($C_{\text{Benzaldehyde}}$, mol cm^{-3}) with time; dependence of benzaldehyde consumption rate (R , $\text{mol mol}^{-1} \text{ h}^{-1}$) with (IA) stirring speed and (IB) mass of catalyst (W , g) for reaction over Au/Al₂O₃. Note: Solid lines provide a guide to aid visual assessment. Reaction conditions: $T=353 \text{ K}$, $P=9 \text{ bar}$

Rate remained constant $> 700 \text{ rpm}$, indicative of minimal gas–liquid/liquid–solid mass transfer contributions to overall rate [45]. Based on these results, the stirring speed was set at 900 rpm for subsequent tests. Rate was invariant over the range of catalyst masses considered in this work, confirming chemical control where external or internal transport constraints do not contribute to the catalytic response [45].

3.2.2 Reaction Mechanism and Solvent Effects

The effect of ring substituents was considered by testing the hydrogenation of substituted benzaldehydes bearing electron donating and withdrawing substituents ($-\text{OCH}_3$, $-\text{CH}_3$, $-\text{CH}_2\text{CN}$, $-\text{CN}$) in the *para* position. In each case, Au/Al₂O₃ showed full selectivity to the corresponding alcohol with decreasing activity in the order: *p*-cyanobenzaldehyde $>$ *p*-formylphenyl acetonitrile $>$ benzaldehyde $>$ *p*-tolualdehyde $>$ *p*-methoxybenzaldehyde. The rate data were subjected to a Hammett correlation in order to probe reaction mechanism [46]. Rate for the substituted benzaldehyde (R_i) is related to benzaldehyde as reference (R) according to

$$\log \left[\frac{R_i}{R} \right] = \rho \times \sigma_p \quad (6)$$

The reaction constant (ρ) provides a measure of the susceptibility to substituent electronic effects while the σ_p factor is an empirical parameter (values taken from [46]) that reflects substituent electron donating/acceptor character [47, 48]. The fit of the experimental rate data is presented in Fig. 3. The low ρ (0.9) extracted from the

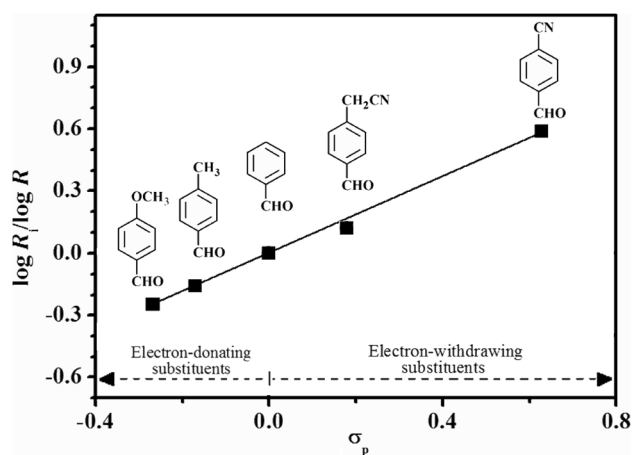


Fig. 3 Hammett plot for the selective $-\text{C}=\text{O}$ group reduction of *para*-substituted benzaldehydes over Au/Al₂O₃ at $T=353 \text{ K}$ and $P=9 \text{ bar}$. Note: Solid line represents fit to Eq. (6)

linear correlation is close to that (0.7) reported for liquid phase hydrogenation of acetophenones over Pd/C [49] and suggests a partial charge in the transition state [7]. A positive reaction constant is consistent with nucleophilic addition [50]. Hydrogen can undergo heterolytic dissociation on the surface of Au/Al₂O₃ to generate H^- that bonds to Au [16]. The hydride ion (H^-) supplied by Au acts as a nucleophile that attacks the positively charged carbon in the $-\text{C}=\text{O}$ group, activated on support Lewis acid sites (Al^{3+}) at the interface with the metal [51]. An electron-donating substituent (in the *para*-position) lowers the reactivity of the carbonyl-carbon which, in turns, lowers the rate of reaction.

The solvent serves to dissolve the reactant and/or products but can also interact with the catalyst and influence performance [45]. Benzaldehyde hydrogenation rate was lower in ethanol ($23 \text{ mol mol}^{-1} \text{ h}^{-1}$) relative to water ($78 \text{ mol mol}^{-1} \text{ h}^{-1}$). The higher rate in aqueous solution can not be due to H_2 solubility as under reaction conditions ($T=353 \text{ K}$, $P=9 \text{ bar}$) solubility is higher in ethanol ($3.7 \mu\text{mol cm}^{-3}$) than water ($0.8 \mu\text{mol cm}^{-3}$) [25]. The dielectric constant (ϵ) provides a measure of solvent capacity to interact with charged surface sites [52]; ϵ for water + ethanol mixtures was calculated following the approach described previously [25]. An increase in rate with solvent dielectric constant (ϵ) is shown in Fig. 4. Competition of solvent and benzaldehyde for adsorption sites [53] can impact on hydrogenation rate where increased surface affinity for the solvent inhibits benzaldehyde activation, resulting in lower reaction rate. Water [54] and alcohols [55] can adsorb on Lewis acid sites through the oxygen of the hydroxyl group and the greater electron density of the hydroxyl oxygen in alcohols must result in a stronger interaction that increases with increasing alcohol chain length [56].

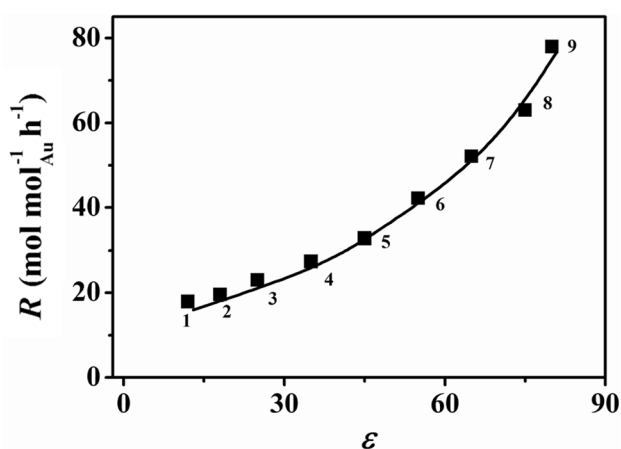


Fig. 4 Benzaldehyde hydrogenation rate (R , $\text{mol mol}^{-1}_{\text{Au}} \text{h}^{-1}$) over $\text{Au}/\text{Al}_2\text{O}_3$ as a function of solvent dielectric constant (ϵ) for reaction in (1) pentanol; (2) butanol; (3) ethanol; ethanol: H_2O =(4) 5:1; (5) 3:1; (6) 1:1; (7) 0.8:1; (8) 0.6:1; (9) H_2O . Note: Solid line provides a guide to aid visual assessment. Reaction conditions: $T=353 \text{ K}$, $P=9 \text{ bar}$

The solvent can also influence reactant activation through hydrogen bonding with the $-\text{C}=\text{O}$ function [28, 29]. This results in charge transfer from the lone pair of the oxygen atom in the carbonyl group to the $\text{O}-\text{H}$ bond in protic solvents [57], the carbonyl-carbon becomes more electrophilic and the rate of reaction increases. Akpa et al. [58] employed DFT calculations to investigate 2-butanone hydrogenation in water over Ru/SiO_2 and concluded that strong interaction between water and 2-butanone lowered the activation energy barrier and enhanced reaction rate. Catalyst (Ru/C) testing has also shown higher 2-butanone hydrogenation rate in solvents with higher ϵ [28]. In contrast, Bertero et al. [27] and Aramendía et al. [30] using Ni/SiO_2 and Pd/AlPO_4 , respectively, reported a decrease in acetophenone hydrogenation rate with increasing ϵ due to solvation that inhibited the adsorption step.

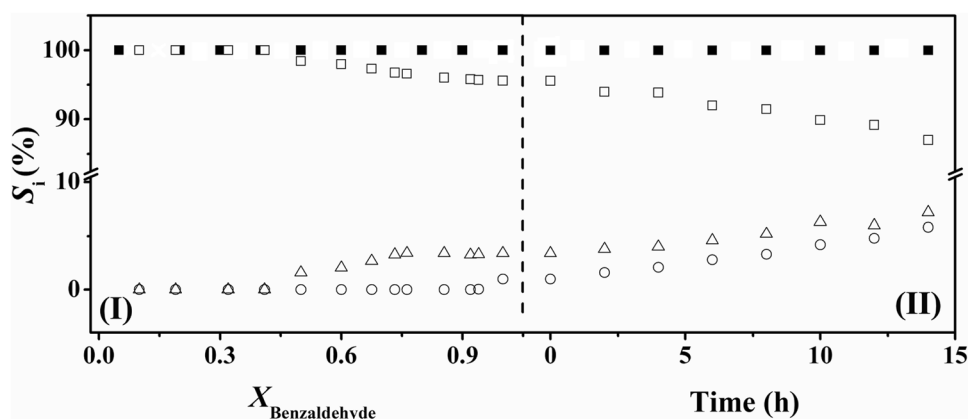
3.2.3 $\text{Au}/\text{Al}_2\text{O}_3$ versus $\text{Pt}/\text{Al}_2\text{O}_3$

The reaction rate (and TOF) recorded for $\text{Pt}/\text{Al}_2\text{O}_3$ was significantly higher (Table 1), which can be attributed to the greater H_2 chemisorption capacity. Reaction over $\text{Au}/\text{Al}_2\text{O}_3$ exhibited full selectivity to the target alcohol at all levels of benzaldehyde conversion (Fig. 5I) to deliver 100% benzyl alcohol yield. This response was probed further by considering benzyl alcohol hydrogenation over $\text{Au}/\text{Al}_2\text{O}_3$; no conversion was detected. The benchmark $\text{Pt}/\text{Al}_2\text{O}_3$ promoted hydrogenolysis to toluene and benzene with a lower yield (95%) to benzyl alcohol at full benzaldehyde conversion. The reaction was prolonged for a further 15 h after complete benzaldehyde conversion (Fig. 5II). Conversion of benzyl alcohol (to toluene and benzene) over $\text{Pt}/\text{Al}_2\text{O}_3$ resulted in a temporal decline in selectivity to the alcohol. This suggests sequential hydrogenation (step I, Scheme 1) and hydrogenolysis (steps II and III) steps. In contrast, full selectivity to benzyl alcohol was maintained with $\text{Au}/\text{Al}_2\text{O}_3$ over extended reaction times with no measurable hydrogenolysis. The results from this work illustrate the potential of supported Au for the selective transformation of substituted benzaldehydes with 100% yield of the target alcohol over prolonged reaction times.

4 Conclusions

The liquid phase ($T=353 \text{ K}$, $P=9.0 \text{ bar}$) hydrogenation of benzaldehyde over (1% w/w) Au (mean size = 3.6 nm) on Al_2O_3 was fully selective with 100% yield of the target benzyl alcohol. Exclusive $-\text{C}=\text{O}$ reduction extended to a range of p -substituted ($-\text{OCH}_3$, $-\text{CH}_3$, $-\text{CH}_2\text{CN}$, $-\text{CN}$) benzaldehydes. The reaction proceeds via a nucleophilic mechanism where electron withdrawing ring substituents elevated rate as demonstrated by the linear Hammett relationship. Selective hydrogenation rate increased with increasing solvent dielectric constant (ϵ), which is accounted for in terms of: (i) competition

Fig. 5 Selectivity (S_i , %) as a function of (I) benzaldehyde fractional conversion ($X_{\text{Benzaldehyde}}$) and (II) time, after 100% conversion of benzaldehyde (in water) had been reached over $\text{Au}/\text{Al}_2\text{O}_3$ (solid symbols) and $\text{Pt}/\text{Al}_2\text{O}_3$ (open symbols); benzyl alcohol (open square, filled square), benzene (open triangle) and toluene (open circle). Reaction conditions: $T=353 \text{ K}$, $P=9 \text{ bar}$



for surface adsorption sites that is more pronounced for alcohols with lower ϵ ; (ii) reactant solvation by solvents with higher ϵ that activates $\text{C}=\text{O}$ for nucleophilic attack. Reaction over a $\text{Pt}/\text{Al}_2\text{O}_3$ benchmark (mean Pt size = 1.7 nm) was non-selective with the formation of by-products from hydrogenolysis. A higher consumption rate recorded for $\text{Pt}/\text{Al}_2\text{O}_3$ can be attributed to greater H_2 chemisorption capacity. Our results demonstrate the potential of Au for ultra-selective aldehyde to alcohol hydrogenation using water as solvent.

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Compliance with Ethical Standards

Conflict of interest The authors declare no conflicts of interest.

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